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Amendment to the Claims:

This listing of claims will replace all prior versions and listing of claims in the application:

Claims 1-14.(Cancelled).

Claim 15 (Previously presented): A compound which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides at least one of:

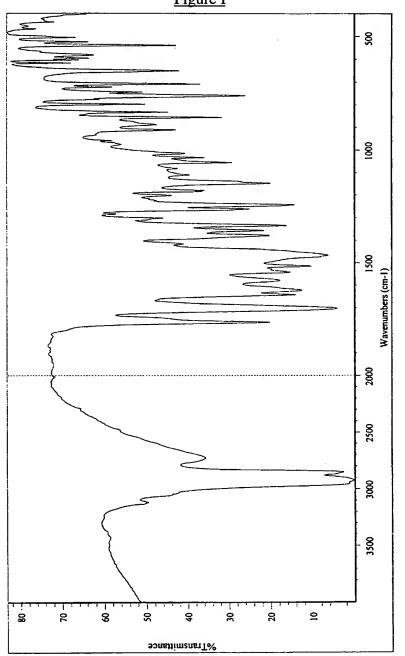
- (i) an infra red spectrum containing peaks at 1763, 912, 856 and 709 cm⁻¹;
- (ii) a Raman spectrum containing peaks at 1762, 1284, 912 and 888 cm⁻¹;
- (iii) a solid-state ¹³C nuclear magnetic resonance spectrum containing peaks at 111.0, 113.6, 119.8, 129.1, 130.9, 131.8, 134.7, 138.7, 146.5, 152.7, 157.5, 169.5. 171.0, 178.7 ppm; and
- (iv) an X-ray powder diffraction pattern having calculated lattice spacings at 5.87, 5.30, 4.69, 4.09, 3.88, 3.61, 3.53 and 3.46 Angstroms.

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Claim 16 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxylbenzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound, in a mineral oil dispersion, provides an infra red spectrum substantially in accordance with Figure I:



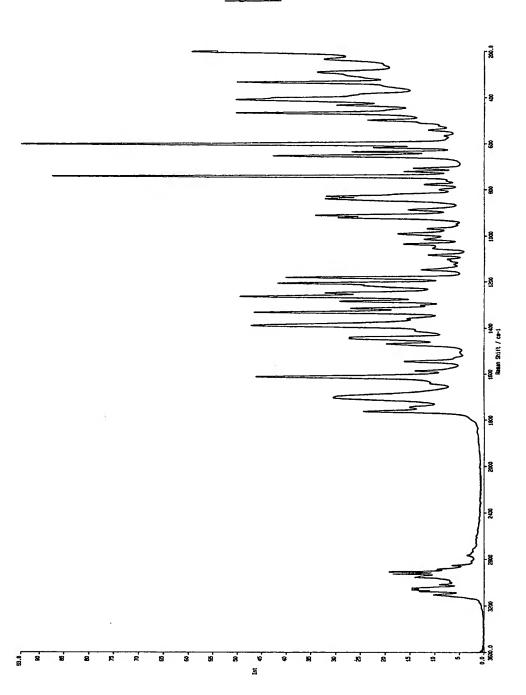


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Claim 17 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides a Raman spectrum substantially in accordance with Figure II:

Figure II



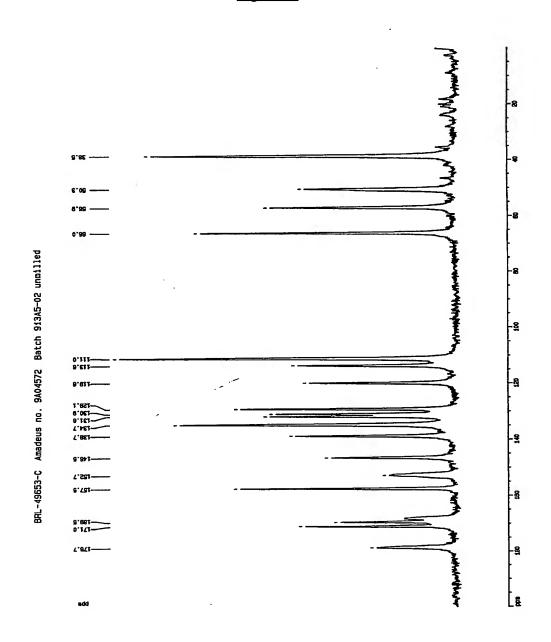
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Claim 18 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides a solid-state ¹³C nuclear magnetic resonance spectrum substantially in accordance with Figure III:

Figure III



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Claim 19 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides a solid-state ¹³C nuclear magnetic resonance spectrum substantially in accordance with Table I:

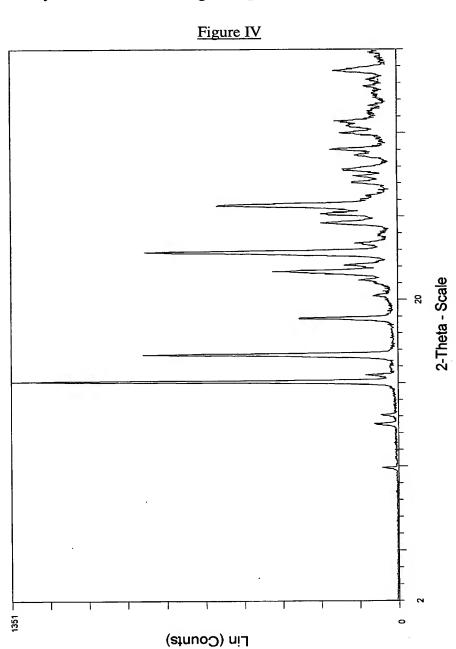
Table I

	Cl	nemical Shift (pp	m)	
	Cı	ionnear omit (pp	,	
38.5	111.0	130.9	146.5	171.0
50.3	113.6	131.8	152.7	178.7
56.9	119.8	134.7	157.5	
66.0	129.1	138.7	169.5	

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Claim 20 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides an X-ray powder diffraction pattern substantially in accordance with Figure IV:



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Claim 21 (Currently Amended): A compound according to claim 15, which provides which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides an X-ray powder diffraction pattern substantially in accordance with Table II:

Table II

Diffraction Angle	Lattice Spacing	
(°2 0)	(Angstroms)	
9.9	8.97	
12.5	7.07	
13.1	6.78	
15.1	5.87	
15.5	5.72	
16.7	5.30	
18.9	4.69	
20.3	4.38	
21.2	4.19	
21.7	4.09	
22.1	4.02	
22.9	3.88	
23.4	3.80	
23.9	3.72	
24.6	3.61	
25.2	3.53	
25.7	3.46	
26.3	3.39	
27.1	3.29	
27.5	3.25	
27.9	3.20	
28.7	3.11	
29.1	3.07	
30.1	2.97	
30.5	2.93	
30.8	2.91	
31.3	2.85	
31.7	2.82	
32.9	2.72	
33.2	2.69	
33.8	2.65	
34.0	2.64	

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Claim 22 (Currently Amended): A compound according to elaim 15 any one of claims 15-21 in isolated form.

Claims 23 and 24 (Cancelled).

Claim 25 (Previously presented): A process for preparing the compound according to claim 15, comprising:

suspending 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl] thiazolidine-2,4-dione, maleic acid salt in acetone; stirring the suspension at an elevated temperature for an extended period of time; and recovering the compound.

Claim 26 (Currently Amended): A process for preparing the compound according to elaim 15 any one of claims 15-21, comprising:

seeding a solution of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl] thiazolidine-2,4-dione, maleic acid salt in denatured ethanol at an elevated temperature with crystals of the compound; cooling the seeded solution; and recovering the compound from the denatured ethanol.

Claim 27 (Currently Amended): A pharmaceutical composition comprising an effective, non-toxic amount of the compound according to elaim 15 any one of claims 15-21 and a pharmaceutically acceptable carrier therefor.

Claim 28 (Currently Amended): A method for the treatment or prophylaxis of diabetes mellitus, conditions associated with diabetes mellitus and certain complications thereof, in a human or non-human mammal which comprises administering an effective, non-toxic, amount of the compound according to claim 15 any one of claims 15-21 to a human or non-human mammal in need thereof.

Claim 29 (Currently Amended): A method for the treatment of Type II diabetes in a human comprising administering an effective, non-toxic amount of the compound according to elaim 15 any one of claims 15-21 to a human in need thereof.

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Claim 30 (Original): A method of claim 29, wherein the administrating comprises oral administration.

Claim 31 (Previously presented): A method of claim 30, wherein the compound is administered in the form of a tablet or capsule for said oral administration.

Claim 32 (Original): A composition according to claim 27 which is adapted for oral administration.

Claim 33 (Original): A composition according to claim 32 which is in the form of a tablet or a capsule for said oral administration.

Claim 34 (Previously presented): A compound which is a polymorphic form of 5-[4-[2-(N-methyl-N-(2-pyridyl)amino)ethoxy]benzyl]thiazolidine-2,4-dione, maleic acid salt, wherein said compound provides:

- (i) an infra red spectrum containing peaks at 1763, 912, 856 and 709 cm⁻¹;
- (ii) a Raman spectrum containing peaks at 1762, 1284, 912 and 888 cm⁻¹;
- (iii) a solid-state ¹³C nuclear magnetic resonance spectrum containing peaks at 111.0, 113.6, 119.8, 129.1, 130.9, 131.8, 134.7, 138.7, 146.5, 152.7, 157.5, 169.5. 171.0, 178.7 ppm; and
- (iv) an X-ray powder diffraction pattern having calculated lattice spacings at 5.87, 5.30, 4.69, 4.09, 3.88, 3.61, 3.53 and 3.46 Angstroms.

Claim 35 (Currently Amended): A pharmaceutical composition consisting essentially of an effective, non-toxic amount of the compound according to elaim 15 any one of claims 15-21 and a pharmaceutically acceptable carrier therefor.

Claims 36-41 (Cancelled).

Claim 42 (New): A method for the preparation of a pharmaceutical composition comprising admixing an effective, non-toxic amount of the compound according to any one of claims 15-21 with a pharmaceutically acceptable carrier therefor.